

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : <b>A61L 27/00, 15/32, 15/40, 31/00</b>		A1	(11) International Publication Number: <b>WO 96/25961</b> (43) International Publication Date: 29 August 1996 (29.08.96)
(21) International Application Number: PCT/GB96/00399 (22) International Filing Date: 22 February 1996 (22.02.96)		(81) Designated States: CA, JP, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  <b>Published</b> <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
(30) Priority Data: 9503492.2 22 February 1995 (22.02.95) GB			
(71) Applicant (for all designated States except US): ED GEISTLICH SÖHNE AG FÜR CHEMISCHE INDUSTRIE [CH/CH]; Bahnhofstrasse 40, CH-6110 Wohlhusen (CH).			
(71) Applicant (for GB only): PETT, Christopher [GB/GB]; Frank B. Dehn & Co., Imperial House, 15-19 Kingsway, London WC2B 6UZ (GB).			
(72) Inventors; and			
(75) Inventors/Applicants (for US only): GEISTLICH, Peter [CH/CH]; Kehrsitenstrasse 19, CH-6362 Stansstad (CH). SPECTOR, Myron [US/US]; Department of Orthopaedic Surgery, Brigham & Women's Hospital, 75 Francis Street, Boston, MA 02115 (US). ECKMAYER, Zdenek [DE/DE]; Rote Turmstrasse 28, D-69469 Weinheim (DE).			
(74) Agents: PETT, Christopher et al.; Frank B. Dehn & Co., 179 Queen Victoria Street, London EC4V 4EL (GB).			
(54) Title: RESORBABLE EXTRACELLULAR MATRIX FOR RECONSTRUCTION OF CARTILAGE TISSUE			
(57) Abstract			
The invention relates to a resorbable extracellular matrix for reconstruction of cartilage tissue comprising predominantly fibres of collagen II.			

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgyzstan	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	KZ	Kazakhstan	SG	Singapore
CH	Switzerland	LJ	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovakia
CM	Cameroon	LR	Liberia	SN	Senegal
CN	China	LT	Lithuania	SZ	Swaziland
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	LV	Latvia	TG	Togo
DE	Germany	MC	Monaco	TJ	Tajikistan
DK	Denmark	MD	Republic of Moldova	TT	Trinidad and Tobago
EE	Estonia	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	UG	Uganda
FI	Finland	MN	Mongolia	US	United States of America
FR	France	MR	Mauritania	UZ	Uzbekistan
GA	Gabon			VN	Viet Nam

- 1 -

## RESORBABLE EXTRACELLULAR MATRIX FOR RECONSTRUCTION OF CARTILAGE TISSUE

5       The present invention concerns an extracellular matrix for reconstruction of cartilage tissue.

In tissue engineering, it has long proved difficult to reconstruct cartilage. Reconstruction of tissue, in general, comprises provision of a matrix which serves as 10 a guide for cells which grow along and between the fibres of the matrix. Hitherto, attempts to reconstruct cartilage, using matrices based on polylactic acid, polyglycolic acid and collagen I or III, required the matrices to be loaded in vitro with chondrocytes prior 15 to implantation of the loaded matrix in an appropriate in vivo site. It had not proved possible simply to implant the matrices of this type at the in vivo site and rely on growth of the native chondrocytes on the 20 surface of the matrix. The need to load the matrix with chondrocytes in vitro prior to implantation gave rise to complications and difficulties in terms of the sterile culture of the chondrocytes.

There is thus a need for a matrix implant for 25 reconstruction of cartilage tissue which will permit ingrowth of native chondrocytes after implantation in vivo. We have now found that these requirements may be met by a matrix of collagen fibres, provided that the collagen is predominantly collagen II.

Collagen occurs in a number of forms in the animal 30 body and different tissues contain different proportions of the respective types. Thus, whereas bone collagen comprises predominantly collagen I and III, cartilage comprises predominantly collagen II together with smaller quantities of collagen VI, IX, X, XI and XIII. 35 Such material differs significantly from collagen sponge material used in medicine and in cosmetics which, being derived from skin and tendons consists of collagen I

- 2 -

and/or III.

According to one aspect of the present invention, therefore, there is provided a resorbable extracellular matrix for reconstruction of cartilage tissue comprising 5 predominantly fibres of collagen II.

As indicated above, such a matrix may contain minor quantities of collagen VI, IX, X, XI and XIII. The matrix according to the invention desirably also contains a hydrogel-like material, for example 10 comprising glycosaminoglycans such as chondroitin sulphate, keratan sulphate, dermatan sulphate and hyaluronic acid, which provides a natural medium in which chondrocytes can become embedded and grow. In general, the matrix according to the invention 15 preferably contains 0.1 to 40% by weight of glycosaminoglycan, for example 5-15% e.g. about 10% by weight.

The matrix according to the invention may either comprise natural cartilage material which has been 20 subjected to defatting and other treatment, leaving the collagen II material together with glycosaminoglycans, or alternatively fibres of purified collagen II may be mixed with glycosaminoglycans and any other required additives. Such additional additives may, for example, 25 include chondronectin or anchorin II to assist attachment of the chondrocytes to the collagen II fibres and growth factors such as cartilage inducing factor (CIF), insulin-like growth factor (IGF) and transforming growth factor  $\beta$  (TGF $\beta$ ).

There exists a wide range of glycosaminoglycans and 30 proteoglycans which have different and sometimes undesirable properties. Thus, although it is possible to incorporate into the collagen matrix glycosaminoglycans from different sources which do not 35 have the same composition, molecular weight and physiological properties as glycosaminoglycans from cartilage, it is particularly preferred to use

- 3 -

glycosaminoglycans from cartilage itself.

It is desirable to subject the collagen matrix to some degree of cross-linking in order to restrict the extent of swelling when the matrix comes in contact with aqueous fluids, while retaining the ability of the matrix to be resorbed. Such swelling leads to loss of strength and shape. However, chemical cross-linking may present physiological disadvantages in terms of pore size which could negatively influence the properties of the collagen. The pore size should optionally be around 0.4 $\mu$  in order to promote chemotaxis and other functions of the cells. The collagen matrix according to the invention may advantageously be manufactured by subjecting cartilage tissue to defatting followed by treatment with a base whereby proteoglycans and glycosaminoglycans are removed.

The cartilage material will normally be that from readily available animal sources such as cattle, sheep or pigs. The preferred material is hyaline cartilage from pigs. This contains the right type of collagen and glycosaminoglycan in desirable proportions and is available in suitably large quantities.

The cartilage is preferably frozen after slaughter and subjected to size reduction, for example to a particle diameter of about 8mm. Before size reduction, the cartilage is preferably soaked in water and mechanically separated from flesh, bone and other unwanted materials.

The particulate cartilage is then preferably subjected to dewatering by treatment with a water miscible organic solvent such as acetone, which also serves to remove some fat. The dewatering shrinks the collagen fibres and separates them from each other so that the subsequent defatting step is optimised. The material is then subjected to defatting with a fat-solvent such as a hydrocarbon eg. hexane, or a halogenated hydrocarbon.

- 4 -

After defatting, the material is thoroughly washed and this is continued until as much water has been taken up as was present originally. By this procedure, the material is optimised for the base-treatment which  
5 follows.

The base-treatment may be effected with a strong alkali, for example and alkali metal hydroxide, eg. sodium hydroxide, for example at a concentration of 1-8% by weight. The treatment time, which will vary  
10 according to the raw material and alkali concentration, is generally 10-48 hours. The treatment temperature will generally be below 20°C. The pH value is normally in the range 12-14. The above conditions are those which are optimal for treatment with NaOH. Treatment  
15 with other bases may require slightly modified conditions.

The base-treatment has the following effects:  
Small quantities of residual fat are saponified.  
The non-collagen, alkali soluble proteins are  
20 denatured, destroyed, dissolved and eliminated.  
The amide groups in the collagen are saponified, thereby changing the electric charge and the isoelectric point of the collagen.

Bacteria, prions and viruses are inactivated and  
25 the collagen is thus sterilised.

It has been found that by this treatment, proteoglycans undergo a useful modification which can be characterised as follows:

the covalent binding of glycosaminoglycans to the  
30 core protein in proteoglycans is cleaved. In this way the glycosaminoglycans can be liberated from the protein of the proteoglycans. This is termed  $\beta$ -elimination.

By the base-treatment, the core protein is split into small peptides which may be removed from the  
35 reaction mixture by dialysis or ultra filtration.

Due to the strong negative charge, the glycosaminoglycans form water soluble salts which can

- 5 -

partially washed from the collagen. These are, however, uncleaved or only slightly cleaved by the base-treatment and can be separated from peptides by dialysis. A part of the glycosaminoglycan (about 3% by weight of the 5 collagen) is bound to the collagen.

Purified glycosaminoglycans may be obtained by dialysis or ultrafiltration of the extract arising from the base-treatment step.

According to the procedure of the present 10 invention, enzymatic treatment is, in general, not used, in view of the variety of different substances present. However, further steps include treating the material with an organic or inorganic acid, such as hydrochloric acid. This has the following effect:

15 Unwanted acid sensitive materials are removed;  
The fibre structure is loosened.

Subsequently, the material is washed, generally until the pH value of the material is between 2.5 and 4.0. The pH value of the material is preferably 20 controlled accurately. The pH value of the material should be uniform across the cross-section of the cartilage.

After the acid treatment, the cartilage is in a 25 water-swelled condition. The material is then subjected to mechanical size-reduction, for example using a colloid mill. The concentration of the collagen in the aqueous medium is then about 2.5-3.5% by weight. The pH value of this mixture should be somewhat acid, for example 3.5-4.5.

30 At this point, glycosaminoglycan may be added to the purified collagen mass, for example in the range 0.1-40% preferably 5 to 15%, of the weight of collagen.

The glycosaminoglycan added to the collagen is 35 preferably that extracted from the natural cartilage, as indicated above. The matrix will then contain, besides collagen II, the glycosaminoglycans hyaluronic acid, chondroitin sulphate and keratan sulphate. The

- 6 -

chondroitin sulphate and keratan sulphate are covalently bonded to the core protein while hyaluronic acid is, indeed, bound to the proteoglycan but not covalently. By the action of the base, the bonding to the core 5 protein is cleaved and the glycosaminoglycan is freed from the protein. Additionally, the core protein is cleaved to small peptides which are readily removed by dialysis or ultrafiltration. It is important that the core protein is removed, since this may be 10 immunologically active. The removal of the core protein is thus an important part of the process of the present invention.

The recovery of the glycosaminoglycans from the base extract may be effected as follows:

15 The medium is neutralised to a pH value in the range 6-8.

The non-collagen proteins are removed by treatment with an adsorbent such as kaolin.

20 Ultrafiltration of the liquid is effected, using a membrane which permits the passage of molecules of weight less than 10000 daltons.

Concentration of the liquid is effected to a solids content of about 2-5 weight percent.

25 After admixture of the glycosaminoglycan with the collagen II, the material is homogenised still further in a colloid mill and the solid content is adjusted to 1.5-2.5 weight percent. This mass can then serve for the production of two types of product, namely a sponge or a collagen sheet.

30 For the production of a sponge, the mass resulting from homogenisation is frozen. The freezing must be precisely controlled, whereby the freezing time, pH value and particle size are exactly maintained in order to provide a reproducible pore size. The frozen product 35 is then freeze-dried. After freeze-drying, the sponge is warmed to 120-140°C for at least 2 hours. In this way, the material is stabilised by light cross-linking.

- 7 -

After the freeze-drying the material is cut to a desired thickness, stamped to the required shape, sterilised and packed.

Because the use of sponges is limited for use in some fields due to insufficient strength, the collagen matrix according to the invention can advantageously be used for the production of collagen sheets, which are suitable for use in a wide range of medical indications.

For the production of collagen sheets, the concentration of purified II collagen fibres in the liquid suspension should be in the range 0.2-3 weight percent, advantageously 0.5-2 weight percent. Air is preferably removed.

A gel is then formed as an intermediate step. The production of the collagen gel can be effected by various techniques known for gel formation.

The gel is then dried, normally on a plate. In this way, not only is water removed but insoluble collagen-glucosaminoglycan products are formed which are very beneficial for the growth of cells.

For either the above products, the matrix according to the invention can be supplemented with active substances. Thus any physiologically active substance which is water soluble or water dispersible can be used. Thus, the matrix may advantageously contain medicinal substances such as antibacterials, eg. taurolidine, or antibiotics such as tetracyclines and gentamycins.

The invention also provides the use of a matrix according to the invention in guided regeneration of cartilage tissue.

The following examples are given by way of illustration only:

Example 1

35

Frozen cartilage from freshly slaughtered pigs was steeped in cold water, thoroughly washed through and

- 8 -

mechanically purified from flesh residues, bones and hard pieces. Subsequently, the material was washed for 30 minutes under flowing water.

Subsequently, the material was ground three times 5 in a homogenizer. The optical particle size at the end of size reduction was about 8mm.

The cartilage pieces were dewatered by washing 4 times with acetone, each time for 8 hours. The cartilage was then defatted by extraction 4 times with 10 n-hexane. Each treatment lasted at least 8 hours. The ratio of hexane to cartilage was 1:10.

After defatting, the cartilage was swelled in drinking water. The ratio of water:material was 10:1. The treatment time was 24 hours.

15 The material was then treated with NaOH (5% by weight) whereby the ratio of cartilage to liquid was 1:4 and the treatment time was 32 hours. During the treatment, the pieces of cartilage were well stirred. Subsequently, the alkali was washed from the cartilage. 20 The original pH of 14 was thereby reduced to 9-11. The dissolved impurities were washed out and separated from the cartilage. The liquid resulting from the alkaline treatment was collected for the recovery of glycosaminoglycan.

25 The collagen material was then treated with strong HCl (about 3% by weight) initially at a pH value under 1.0. The treatment time was 4-6 hours.

Subsequently, the material was washed with cold water long enough for the pH value to rise to 3-3.5. 30 All impurities were removed and the product was a salt-free collagen mass, suitable for production of a sponge or other collagen material. For that purpose, the cartilage mass may be, according to the intended result, degassed, frozen and freeze-dried.

- 9 -

Example 2

The extract resulting from alkaline treatment in Example 1 contained glycosaminoglycan, alkali, denatured 5 proteins and salts. The extract was firstly neutralised with HCl, the pH value after neutralisation being 6. The extract was then treated with a filter aid, namely kieselguhr, which had the effect of removing the denatured proteins. 0.5 weight percent of kieselguhr 10 was introduced into the extract and removed by filtration together with the denatured protein.

The supernatant was then submitted to ultrafiltration using a membrane having a molecular weight cut off at about 10000 daltons. In this way, 15 salts were removed to leave purified glycosaminoglycan.

The glycosaminoglycan solution so obtained was admixed with collagen material from above to provide a collagen II matrix containing glycosaminoglycan.

20 Example 3

(1) Determination of hexosamine and amino acid residues in collagen sponges and fleeces

25 Each sample, exactly weighed (about 10 mg) was hydrolysed in 10 ml of 3M or 6M HCl at 1.05°C for 15 or 20 hours under purified nitrogen in a sealed tube. After cooling the tube in a refrigerator and opening the tube, the contents were transferred to a 25 ml long neck 30 flask and dried at 40°C in a vacuum-rotation dryer (Rotavapor RE120, Büchi, Switzerland) under water jet vacuum. After dissolving the residue in 5ml water, the residue was again dried under water jet vacuum. Subsequently, the residue was taken up in 5ml loading 35 buffer (0.2M relative to Na<sup>+</sup>) at pH 2.2. For determination of the glucosamine and galactosamine values, after previous dilution of an aliquot with

- 10 -

loading buffer (1+10) 150 µl of the sample hydrolysed in 3M HCl was injected into the cartouche of an amino acid analyser (AlphaPlus, type 4151, Pharmacia-LKB, Freiburg) and evaluated by comparison with a standard with the help of a computer (Shimadzu, Dusseldorf). The same procedure was effected with the sample hydrolysed in 6M HCl, wherein 50 µl were injected in a further test cartouche. The double hydrolysis in 3M and 6M HCl is necessary for optimisation of the hexosamine and amino acid analysis since the maximal values for hexosamine and also tyrosine are only obtained after hydrolysis in 3M HCl while maximal values are only obtained for valine, isoleucine and leucine after hydrolysis in 6M HCl.

15

2. Determination of native collagen content in collagen sponges and fleeces

20 25-30 mg (exactly weighed out) of sample were introduced into 30 ml 0.1M sodium hydrogen carbonate solution (pA, Merck, Darmstadt) pH 8.2 to which 1.5 ml of a 6 mg/ml trypsin solution (lyophilised preparation from bovine pancreas, Boehringer, Mannheim) and incubated for 8 hours at 23±1°C in a shaking water bath (Julabo SWI, Seelbach). After cooling the sample in a cold room to 4°C, it was centrifuged at 4°C in a 60 Ti-Rotor (Beckman, Munich) at 32000 RpM for 30 minutes. The residue was filtered in a stirred ultra filtration cell (Mod 8010, Amicon, Witten) through a Diaflow-Filter 30 PM 10 (Amicon, Witten) of diameter 25 mm and 1 ml of the filtrate was hydrolysed in 6M HCl for 20 hours at 105°C. The further working up and analysis of the hydrolysate is identical with that described under (1) above with the exception that the further uptake of the sample after twice evaporating to dryness, was in 150 µl loading buffer, whereby 150 µl was injected into the test cartouche of the amino acid analyser. The

- 11 -

hydroxyproline value obtained after the amino acid analysis (in  $\mu\text{mol/g}$  starting substance), represents the part of the degradable collagen in the sample. When the hydroxyproline value of a parallel hydrolysis (6M HCl) and analysed sample (see (1) above) which represents the total collagen content, is compared with the hydroxyproline value, the percentage proportion of the "native", that is trypsin non-degradable collagen is indicated.

The results are shown in the following table.

Table

		<u><math>\mu\text{mol/g}</math></u>	<u><math>\text{mol}/1000 \text{ mol}</math></u>
15	Hydroxyproline	795.4	97
	Aspartic acid	381.7	47
	Threonine	190.1	23
	Serine	257.0	31
	Glutamic acid	691.3	84
20	Proline	913.2	112
	Glycine	2614.6	320
	Alanine	864.9	106
	Cysteine/2	11.5	2
	Valine	195.7	24
25	Methionine	62.7	8
	Isoleucine	92.8	11
	Leucine	229.9	28
	Tyrosine	27.0	3
	Phenylalanine	119.9	15
30	Histidine	39.8	5
	Hydroxylysine	126.4	15
	Lysine	173.5	21
	Arginine	395.5	48
35	<b>Total</b>	<b>8182.9</b>	<b>1000</b>

- 12 -

	Glucosamine	9.68	1.18
	Galactosamine	46.30	5.66
	Total Hydroxyproline	795.4 $\mu\text{mol/g}$	
5	Trypsin-degradable hydroxyproline	36.9 $\mu\text{mol/g}$	
	"Native" collagen content	95.4 %	

- 13 -

Claims

1. A resorbable extracellular matrix for reconstruction of cartilage tissue comprising predominantly fibres of collagen II.
- 5
2. A matrix as claimed in claim 1 further comprising from 0.1 to 40% by weight, the example 5 to 15% by weight, of a glycosaminoglycan.
- 10
3. A matrix as claimed in claim 2 wherein the glycosaminoglycan is chondroitin sulphate, keratan sulphate, dermatan sulphate or hyaluronic acid.
- 15
4. A matrix as claimed in any one of claims 1 to 3 further comprising chondronectin and/or anchorin II.
- 20
5. A matrix as claimed in any one of claims 1 to 4 in which the collagen is cross-linked without becoming non-resorbable.
- 25
6. A matrix as claimed in any preceding claim which is derived from natural cartilage.
7. A matrix as claimed in claim 6 derived from cattle, sheep or pigs.
- 30
8. A matrix as claimed in claim 7 which is derived from hyaline cartilage from pigs.
9. A process for preparing a matrix as claimed in claim 1 in which cartilage tissue is subjected to defatting, followed by treatment with a base.
- 35
10. A process as claimed in claim 9 in which a glycosamine is impregnated into the matrix.

- 14 -

11. Use of a matrix as claimed in any one of claims 1 to 8 in the manufacture of a guided tissue regeneration implant.

5 12. A method of regenerating cartilage tissue in the human or non-human animal body comprising implantation of a matrix as claimed in any one of claims 1 to 8.

## INTERNATIONAL SEARCH REPORT

International Application No

PL./GB 96/00399

A. CLASSIFICATION OF SUBJECT MATTER  
 IPC 6 A61L27/00 A61L15/32 A61L15/40 A61L31/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
 IPC 6 A61L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO,A,93 11723 (REGEN CORP) 24 June 1993 see examples	1-12
X	WO,A,93 19168 (MOUNT SINAI HOSPITAL CORP) 30 September 1993 see claims	1-3
Y	WO,A,90 13302 (BRIGHAM & WOMENS HOSPITAL) 15 November 1990 see page 17 - page 18; table 1	1-12
P,Y	WO,A,95 18638 (GEISTLICH SOEHNE AG ;HOLMES MICHAEL JOHN (GB); GEISTLICH PETER (CH) 13 July 1995 see the whole document	1-12
	---	-/-

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

## \* Special categories of cited documents :

- \*'A' document defining the general state of the art which is not considered to be of particular relevance
- \*'E' earlier document but published on or after the international filing date
- \*'L' document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*'O' document referring to an oral disclosure, use, exhibition or other means
- \*'P' document published prior to the international filing date but later than the priority date claimed

- \*'T' later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

- \*'X' document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

- \*'Y' document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

- \*'A' document member of the same patent family

1

Date of the actual completion of the international search	Date of mailing of the international search report
24 June 1996	08.07.96
Name and mailing address of the ISA  European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 Rijswijk Tel. (+ 31-70) 340-2040, Tx. 31 651 epo nl. Fax (+ 31-70) 340-3016	Authorized officer  ESPINOSA, M

## INTERNATIONAL SEARCH REPORT

International Application No PL./GB 96/00399
---

## C/(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO,A,93 10722 (RES DEV FOUNDATION) 10 June 1993 see claims ---	1
A	FR,A,2 679 778 (COLETICA) 5 February 1993 see claims ---	1
A	WO,A,90 05755 (COLLAGEN CORP) 31 May 1990 see examples 1-7 ---	1
A	WO,A,92 13565 (SHAW ROBERT F) 20 August 1992 see claims; examples -----	1

1

**INTERNATIONAL SEARCH REPORT**

International application No.

PCT/GB96/00399

**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.:

because they relate to subject matter not required to be searched by this Authority, namely:

Remark: Although claim 12 is directed to a method of treatment of the human or animal body the search has been carried out and based on the alleged effects of the product.

2.  Claims Nos.:

because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3.  Claims Nos.:

because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

**Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)**

This International Searching Authority found multiple inventions in this international application, as follows:

1.  As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2.  As all searchable claims could be searched without effort justifying an additional fee, thus Authority did not invite payment of any additional fee.

3.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4.  No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

**Remark on Protest**

The additional search fees were accompanied by the applicant's protest.



No protest accompanied the payment of additional search fees.

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/GB 96/00399

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO-A-9311723	24-06-93	US-A-	5306311	26-04-94
		AU-B-	3229193	19-07-93
		CA-A-	2125967	24-06-93
		EP-A-	0617598	05-10-94
		JP-T-	7505792	29-06-95
-----	-----	-----	-----	-----
WO-A-9319168	30-09-93	US-A-	5326357	05-07-94
		AU-B-	3881693	21-10-93
		CA-A-	2131904	30-09-93
		EP-A-	0631619	04-01-95
		JP-T-	7505620	22-06-95
-----	-----	-----	-----	-----
WO-A-9013302	15-11-90	AU-B-	5654990	29-11-90
-----	-----	-----	-----	-----
WO-A-9518638	13-07-95	NONE		
-----	-----	-----	-----	-----
WO-A-9310722	10-06-93	AU-B-	663150	28-09-95
		AU-B-	3224593	28-06-93
		CA-A-	2124190	10-06-93
		CN-A-	1079912	29-12-93
		EP-A-	0614345	14-09-94
		FI-A-	942425	25-05-94
		JP-T-	7501465	16-02-95
		NO-A-	941917	24-05-94
		NZ-A-	245286	27-06-95
		ZA-A-	9209119	21-09-94
-----	-----	-----	-----	-----
FR-A-2679778	05-02-93	AU-B-	660045	08-06-95
		AU-B-	2474592	02-03-93
		EP-A-	0641225	08-03-95
		FI-A-	940472	01-02-94
		WO-A-	9302718	18-02-93
		JP-T-	7509143	12-10-95
		NO-A-	940279	30-03-94
-----	-----	-----	-----	-----
WO-A-9005755	31-05-90	US-A-	5162430	10-11-92
		AU-B-	638687	08-07-93
		AU-B-	4660989	12-06-90
		CA-A-	2003538	21-05-90
		EP-A-	0444157	04-09-91

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/GB 96/00399

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO-A-9005755		JP-T- 4502027	09-04-92	
		US-A- 5306500	26-04-94	
		US-A- 5510418	23-04-96	
		US-A- 5376375	27-12-94	
		US-A- 5413791	09-05-95	
		US-A- 5475052	12-12-95	
		US-A- 5523348	04-06-96	
		US-A- 5446091	29-08-95	
		US-A- 5470911	28-11-95	
		US-A- 5476666	19-12-95	
		US-A- 5510121	23-04-96	
		US-A- 5324775	28-06-94	
		US-A- 5328955	12-07-94	
		US-A- 5264214	23-11-93	
		US-A- 5308889	03-05-94	
		US-A- 5292802	08-03-94	
		US-A- 5304595	19-04-94	
-----	-----	-----	-----	-----
WO-A-9213565	20-08-92	US-A- 5206023	27-04-93	
		AT-T- 121943	15-05-95	
		AU-B- 667032	07-03-96	
		AU-B- 1412892	07-09-92	
		CA-A- 2101556	01-08-92	
		CN-A- 1064813	30-09-92	
		DE-D- 69202332	08-06-95	
		DE-T- 69202332	04-01-96	
		EP-A- 0569541	18-11-93	
		ES-T- 2072144	01-07-95	
		JP-T- 6505258	16-06-94	
		US-A- 5368858	29-11-94	
-----	-----	-----	-----	-----